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Kautzky

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- (54) **METHOD FOR ANALYZING HEMOSTASIS**
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- (*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 1140 days.

This patent is subject to a terminal disclaimer.

5,096,669	A	3/1992	Lauks et al.	
5,100,805	A	3/1992	Ziege et al.	
5,114,860	A	5/1992	Hayashi	
5,167,145	A	12/1992	Butler et al.	
D337,164	S	7/1993	Lauks et al.	
5,416,026	A	5/1995	Davis	
5,447,440	A	9/1995	Davis et al.	
5,534,226	A	7/1996	Gavin et al.	
5,580,744	A	12/1996	Zweig	
5,628,961	A	5/1997	Davis et al.	
6,060,323	A	5/2000	Jina	
6,066,504	A	5/2000	Jina	
6,136,271	A	10/2000	Lorincz et al.	
6,196,979	B1	3/2001	Virtanen	
6,200,532	B1	3/2001	Wu et al.	
6,327,031	B1	12/2001	Gordon	
6,338,821	B1	1/2002	Jina	
6,438,498	B1	8/2002	Opalsky et al.	
6,524,861	B1	2/2003	Anderson	
6,787,363	B2 *	9/2004	Cohen et al.	436/69
6,797,519	B2 *	9/2004	Cohen et al.	436/69
7,261,861	B2 *	8/2007	Kautzky	422/73

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G01N 37/00 (2006.01)
(52) **U.S. Cl.** **436/69**; 73/64.41; 422/73
(58) **Field of Classification Search** 436/69;
73/64.41; 422/73
See application file for complete search history.

(56) **References Cited**
U.S. PATENT DOCUMENTS
3,854,324 A 12/1974 Altshuler et al.
4,149,405 A 4/1979 Ringrose
4,202,204 A 5/1980 Hartert
4,341,111 A 7/1982 Husar
4,695,956 A 9/1987 LeVeen et al.
5,075,077 A 12/1991 Durley, III et al.

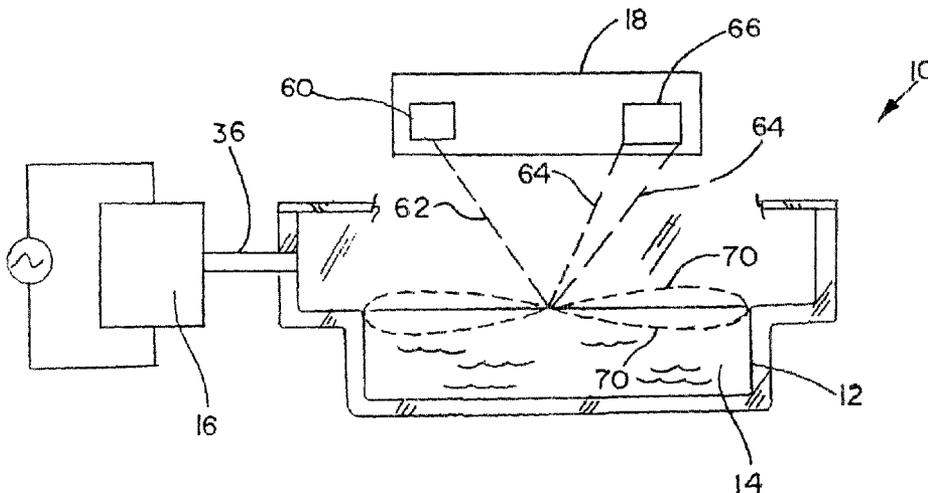
FOREIGN PATENT DOCUMENTS
WO WO-01/96879 A2 12/2001

OTHER PUBLICATIONS
Voleisis et al., "Ultrasonic method for the whole blood coagulation analysis," *Ultrasonics*, vol. 40, May 2002, pp. 101-107.
* cited by examiner

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(57) **ABSTRACT**
A method and device for blood hemostasis analysis is disclosed. A blood sample is displaced to reach a resonant state. The resonant frequency of the blood sample is determined before, during and after a hemostasis process. The changes in the resonant frequency of the blood sample are indicative of the hemostasis characteristics of the blood sample.

13 Claims, 8 Drawing Sheets



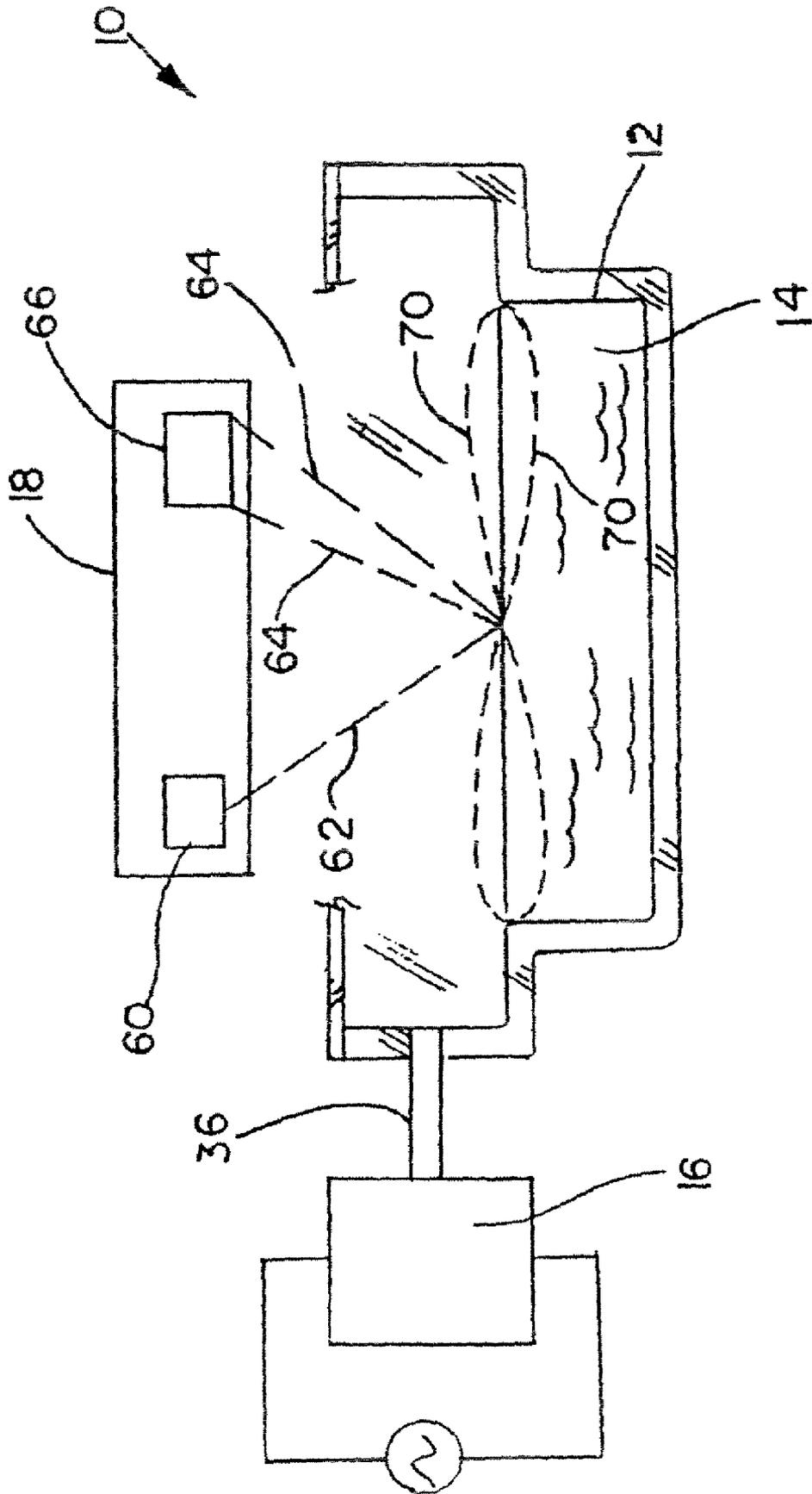


FIG. 1

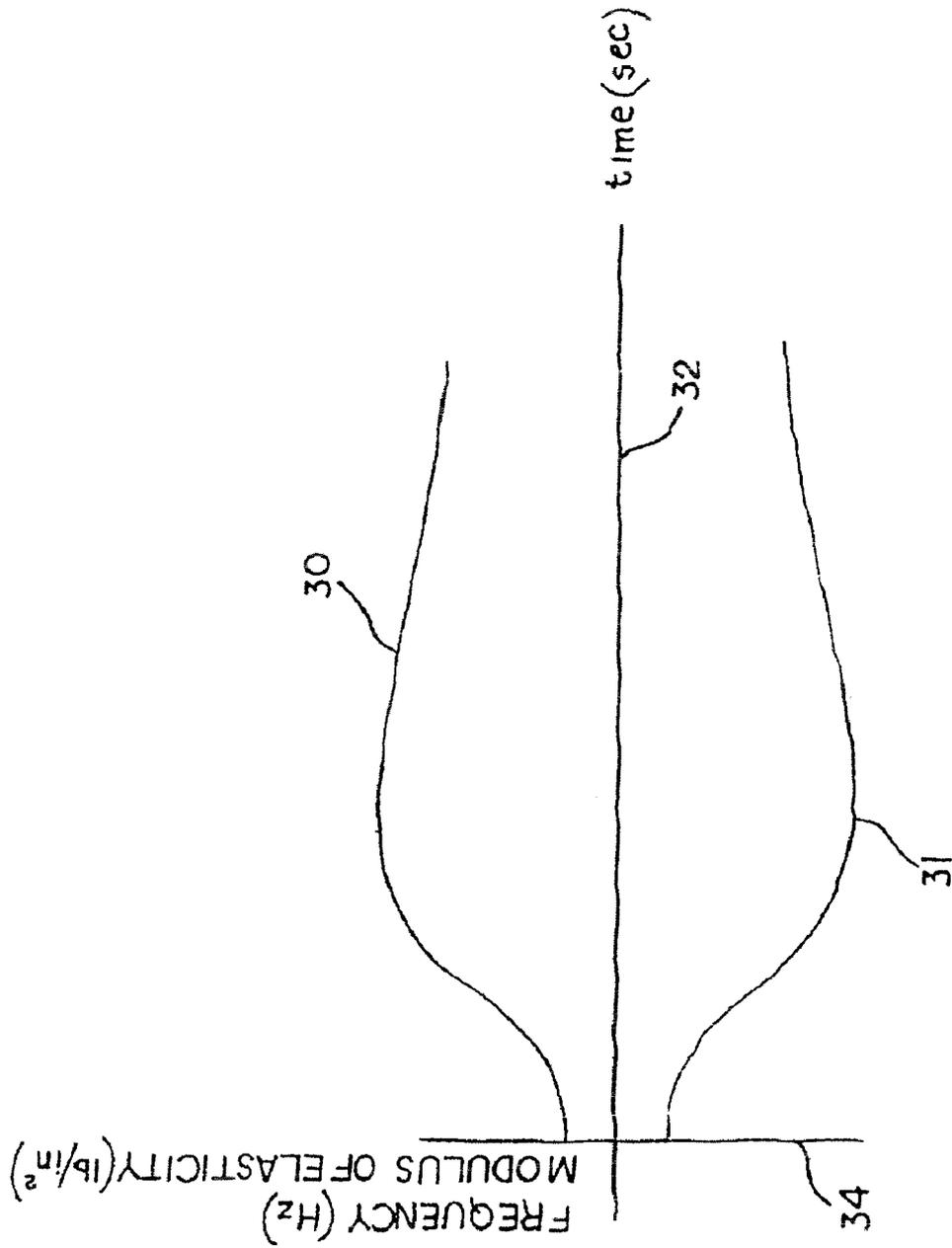


FIG. 2

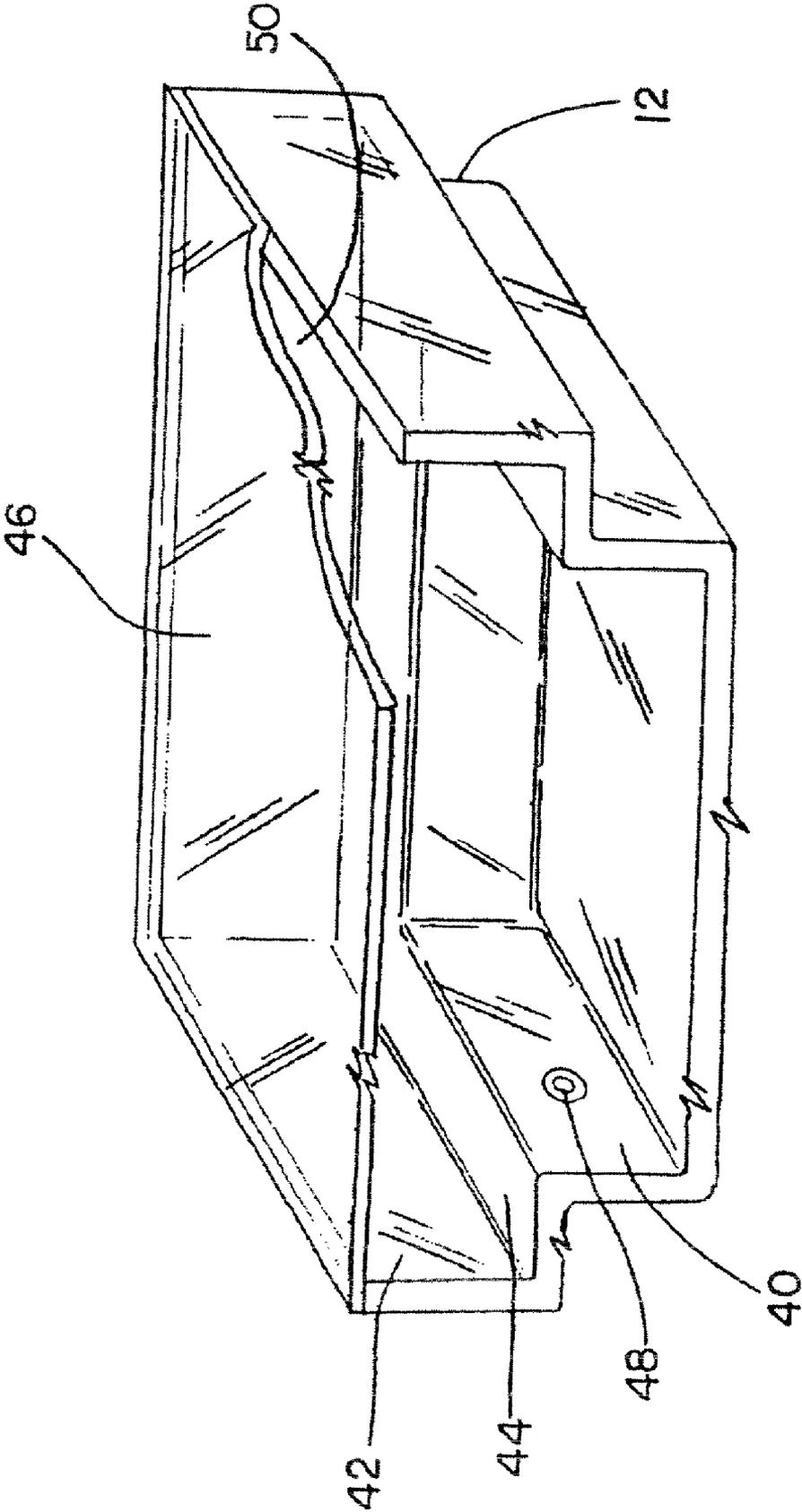


FIG. 3

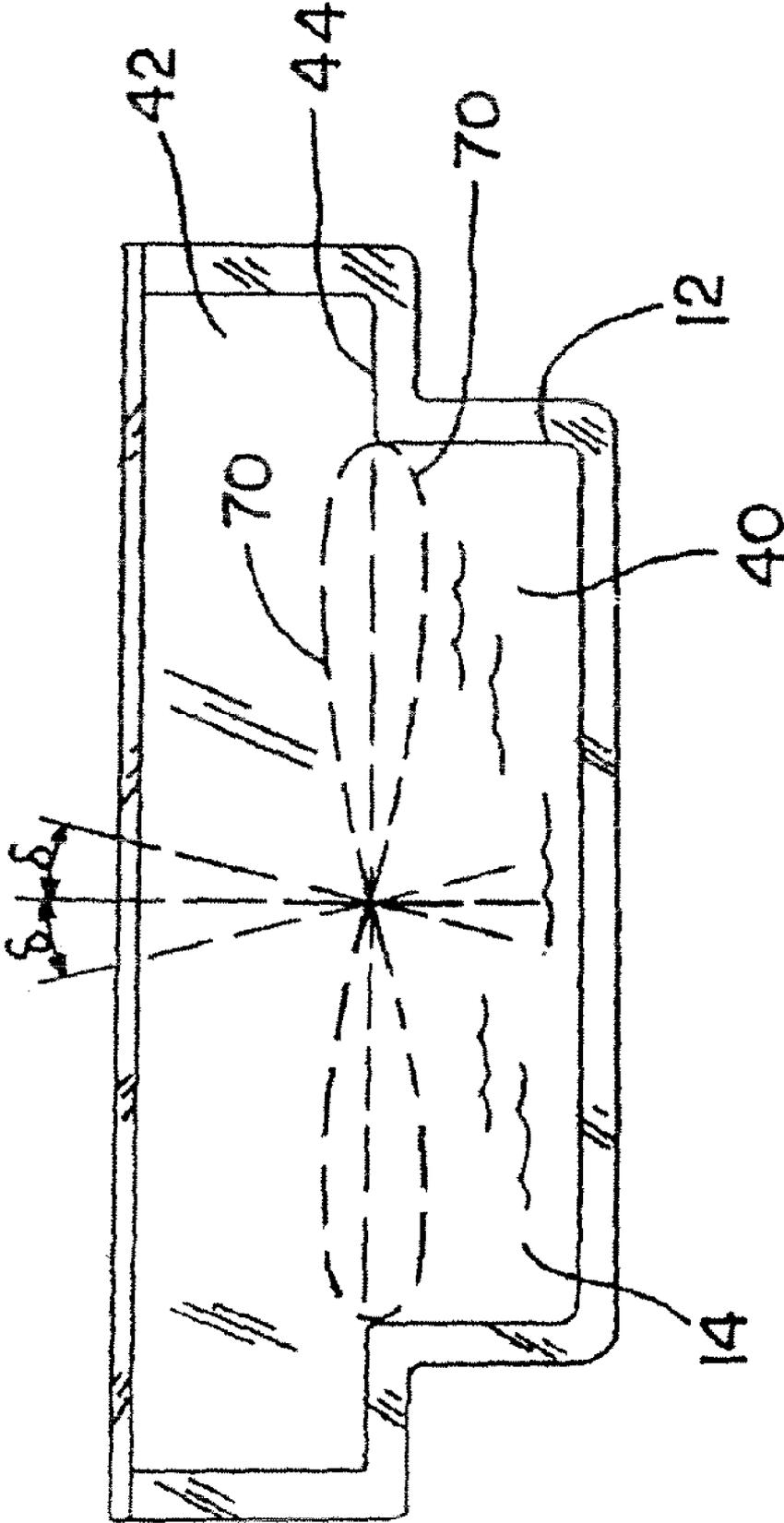


FIG. 4

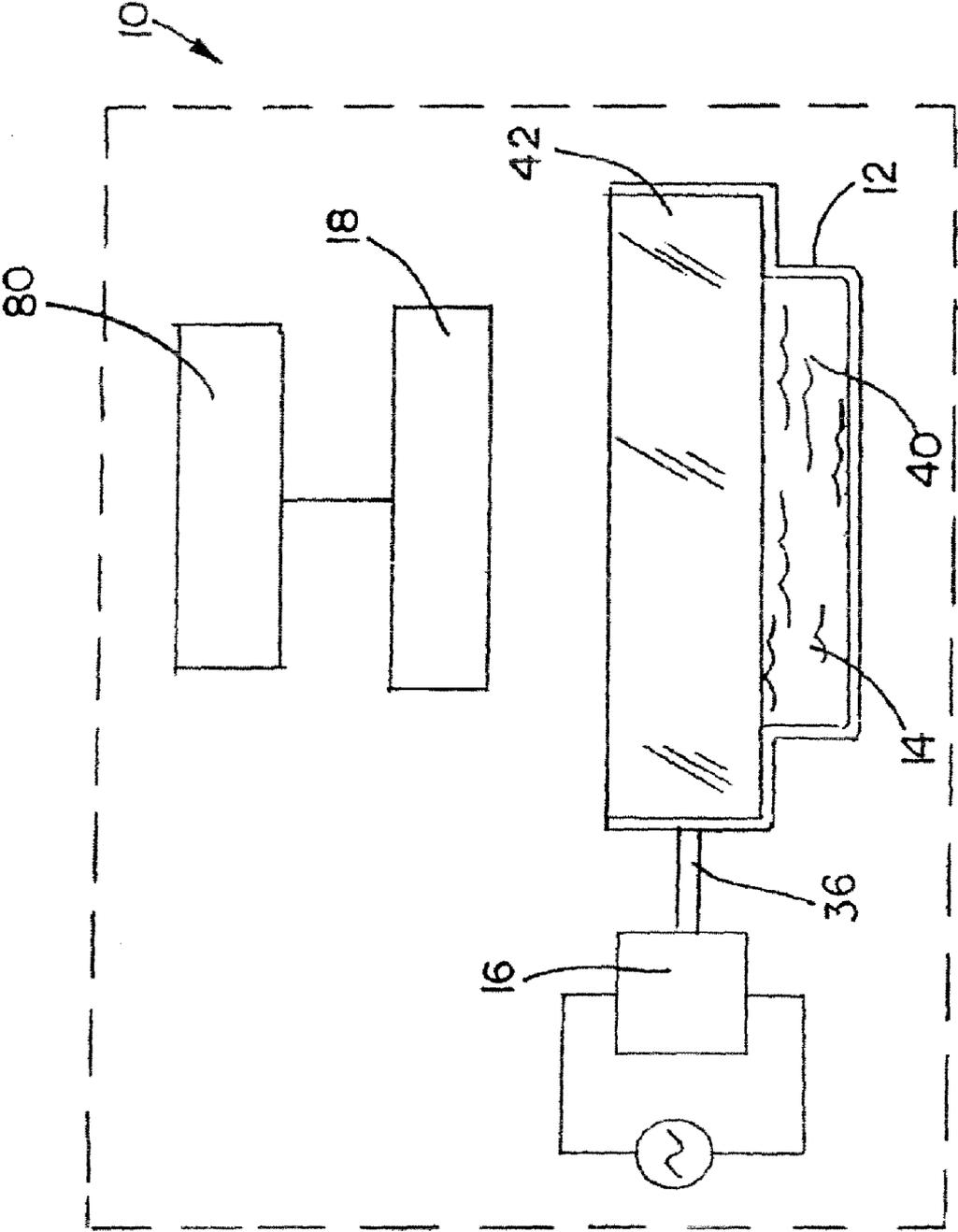


FIG. 5

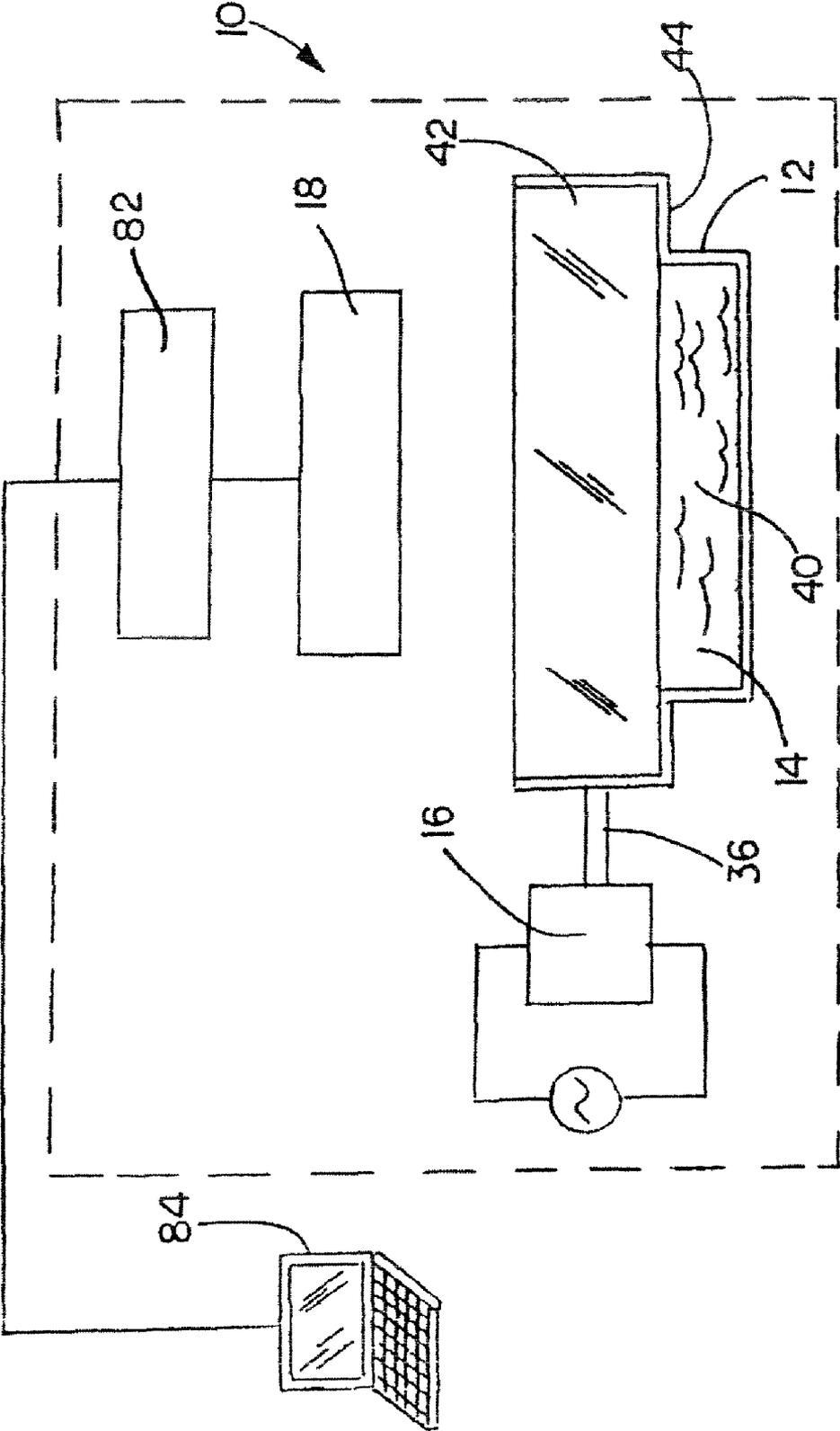


FIG. 6

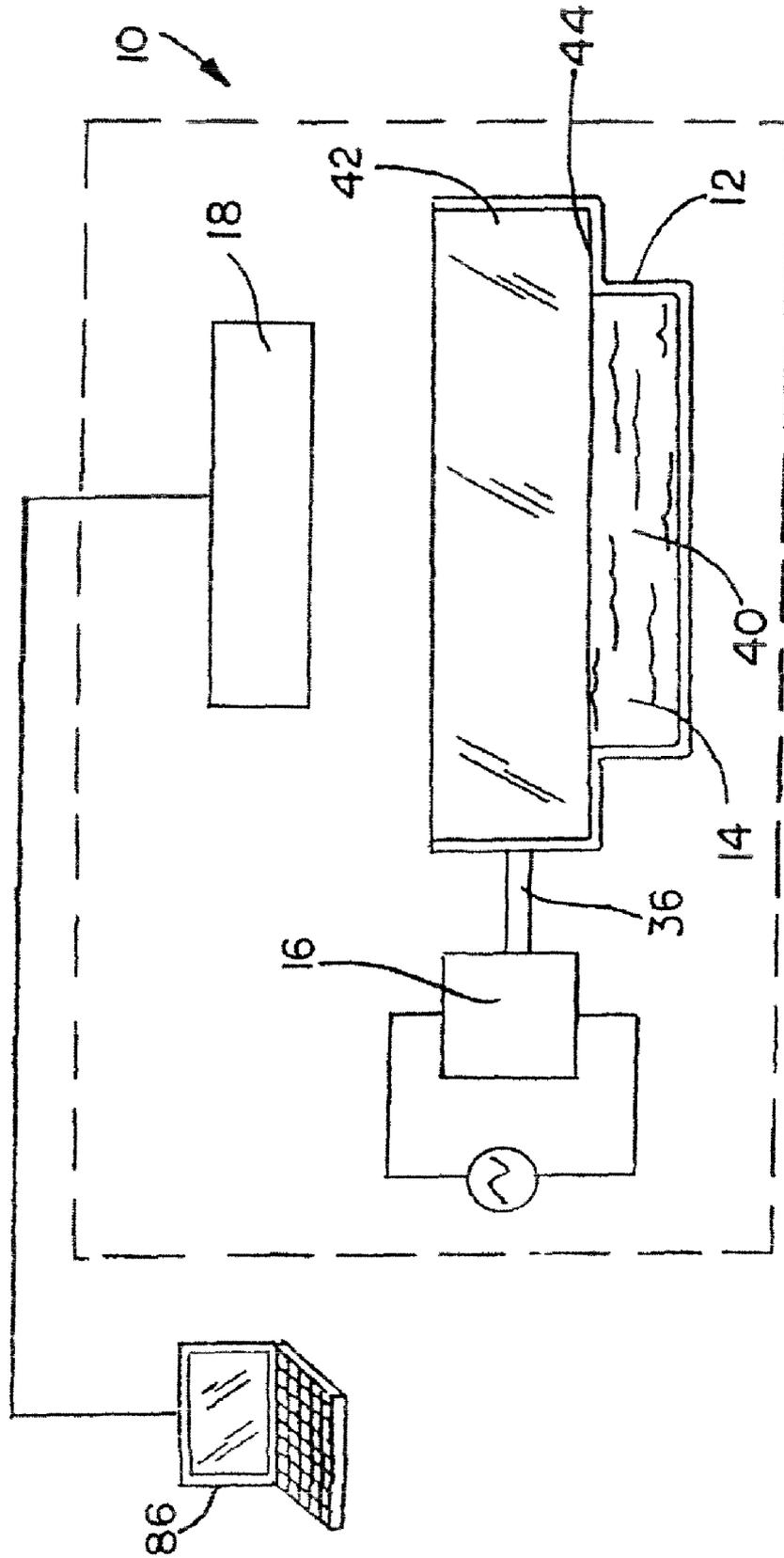


FIG. 7

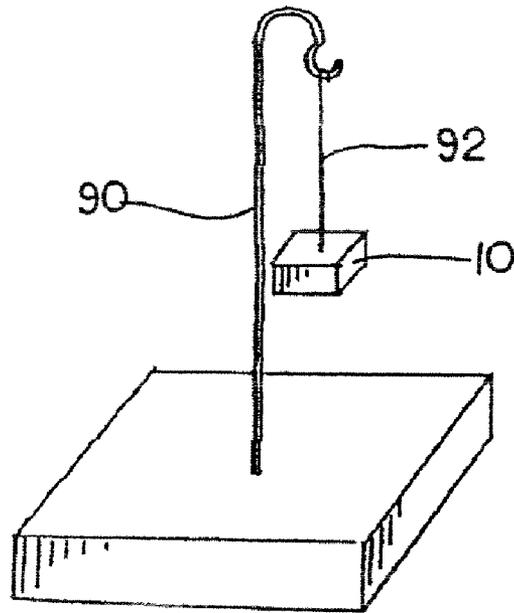


FIG. 8

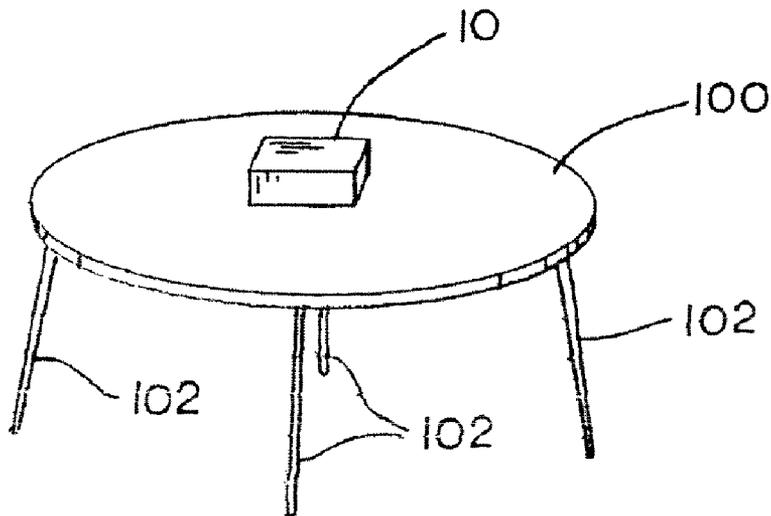


FIG. 9

METHOD FOR ANALYZING HEMOSTASIS

CROSS-REFERENCE TO RELATED APPLICATIONS

This patent is a divisional of U.S. patent application Ser. No. 10/442,229 filed Apr. 24, 2003, the disclosure of which is hereby expressly incorporated for all purposes.

FIELD OF THE DISCLOSURE

This disclosure relates generally to blood analysis, and more particularly, to a blood hemostasis analyzer and method.

BACKGROUND

Blood is in liquid form when traveling undisturbed in bodily passageways. However, an injury may cause rapid clotting of the blood at the site of the injury to initially stop the bleeding, and thereafter, to help in the healing process. An accurate measurement of the ability of a patient's blood to coagulate in a timely and effective fashion and to subsequent lysis is crucial to certain surgical and medical procedures. Also, accurate detection of abnormal hemostasis is of particular importance with respect to appropriate treatment to be given to patients suffering from clotting disorders.

Blood hemostasis is a result of highly complex biochemical processes that transform the blood from a liquid state to a solid state. Characteristics of blood, such as strength of the clot, infer that the mechanical properties of the blood are important in determining characteristics rather than the viscosity of the blood when in a liquid state.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a schematic diagram of a blood hemostasis analyzer constructed in accordance with the teachings of the instant disclosure.

FIG. 2 is a graph representing hemostasis characteristics of a blood sample in accordance with the teachings of the instant disclosure.

FIG. 3 is a perspective and exploded sectional view of a container for holding a blood sample in accordance with the teachings of the instant disclosure.

FIG. 4 is a schematic view of the container of FIG. 3 having therein a blood sample and vibrating the blood sample in accordance with the teachings of the instant disclosure.

FIG. 5 is a schematic view of an analyzer in accordance with the teachings of the instant disclosure.

FIG. 6 is a schematic view of an analyzer in accordance with the teachings of the instant disclosure.

FIG. 7 is a schematic view of an analyzer in accordance with the teachings of the instant disclosure.

FIG. 8 is a perspective view of a first exemplary stand for a blood hemostasis analyzer constructed in accordance with the teachings of the instant disclosure.

FIG. 9 is a perspective view of a second exemplary stand for a blood hemostasis analyzer constructed in accordance with the teachings of the instant disclosure.

DETAILED DESCRIPTION

Referring to FIG. 1, a blood hemostasis analyzer 10 in accordance with the teachings of the instant disclosure is generally shown. The analyzer 10 operates under the principle that because hemostasis of a blood sample changes the

blood sample from a liquid state to a gel-like state, and the modulus of elasticity of the blood sample controls the natural frequency of the blood sample, measuring the changes in the natural frequency of the blood sample during hemostasis provides the hemostasis characteristics of the blood sample. In keeping with this principle, the disclosed blood hemostasis analyzer 10 measures the changes in the fundamental natural frequency of a blood sample during hemostasis and lysis processes to provide hemostasis characteristics of the blood sample. To practice the foregoing principle, the analyzer 10 generally includes a container 12 for holding a blood sample 14, a shaker 16 for displacing the container 12 to thereby excite the blood sample 14 to a resonant vibration, and a sensor 18 for measuring the resulting amplitude of the blood sample 14.

An exemplary method by which the disclosed blood hemostasis analysis is performed will now be described. Vibration of a liquid at resonance closely resembles sloshing, which is analogous to the motion of a pendulum. Accordingly, as blood transitions from a liquid state to a gel-like state and possibly to a solid state during clotting, the fundamental natural frequency of the blood increases. The disclosed exemplary method measures the changes in the fundamental natural frequency of the blood sample 14 during hemostasis/clotting and lysis processes.

Initially a blood sample 14 is placed in the container 12. The container 12 is then vibrated by the shaker 16 so that the blood sample 14, which is initially in a liquid state, is vibrating in a linear sloshing mode. A liquid typically vibrates near its first fundamental natural frequency in a sloshing mode, which can be defined as the swinging of the entire mass of the liquid in a container, hence the analogy to a pendulum. The amplitude of the sloshing reaches maximum when the blood sample 14 is vibrated at its fundamental natural frequency. Thus, to initially excite the blood sample 14 to resonance, the shaker 16 vibrates the container 12 at or very near the fundamental natural frequency of the blood sample 14. Furthermore, the shaker 16 vibrates the container 12 at or very near the fundamental natural frequency of the blood sample 14 as this frequency changes throughout the hemostasis and possibly lysis processes.

One of ordinary skill in the art will readily appreciate the numerous methods by which the shaker 16 can vibrate the container 12 at or near the fundamental natural frequency of the blood sample 14 throughout the hemostasis and lysis processes. However, in the disclosed example, the container 12 is initially vibrated at a frequency below the fundamental natural frequency of the blood sample 14. The frequency is then increased in small steps and concurrently, the resulting displacement amplitudes of the blood sample 14 are measured. As the frequency of vibration of the container 12 increases to near the blood sample's fundamental natural frequency, the displacement amplitude of the blood sample 14 will dramatically increase. The displacement amplitude of the blood sample 14 will reach maximum at its fundamental natural frequency. Thus, monitoring the displacement amplitude of the blood sample 14 for a maximum provides a value for the fundamental natural frequency of the blood sample 14 when that maximum is reached.

As the hemostasis process continues, the foregoing method of finding the fundamental natural frequency of the blood sample 14 is repeated. The measured fundamental natural frequencies of the blood sample 14 when plotted vs. time result in a curve 30 similar to that shown in FIG. 2. Curve 30 is typically represented with its mirror image relative to the x-axis, which is shown as curve 31. The shape of the curve 30 is indicative of blood hemostasis characteristics. The x-axis

32 represents time, while the y-axis 34 represents the fundamental natural frequency of the blood sample 14 during the hemostasis and lysis processes. One of ordinary skill in the art will appreciate that since frequency of the blood sample 14 is proportional to the modulus of elasticity of the blood sample 14, the y-axis also represents the changes in the modulus of elasticity of the blood sample 14 during hemostasis and lysis processes.

One of ordinary skill in the art will readily appreciate that the size of the frequency step by which the vibration frequency of the container 12 is increased or decreased during testing will affect how quickly and efficiently the fundamental natural frequency of the blood sample 14 is pinpointed. For instance, a very large frequency step may not provide a detailed frequency resolution to locate a near accurate measure of the fundamental natural frequency of the blood sample 14. On the other hand, a very small frequency step may not provide a rapid approach to pinpointing the fundamental natural frequency of the blood sample 14. Accordingly, in order to find the fundamental natural frequency of the blood sample within the frequency range by which the container 12 is vibrated, it may be necessary to search for the fundamental natural frequency of the blood sample 14 by changing the frequency step and/or adding or subtracting the frequency step from the vibration frequency of the container 12 in a methodical manner. Numerous mathematical algorithms and methods are well known to those of ordinary skill in the art, by which the frequency step can be methodically varied to provide a rapid pinpointing of a peak in amplitude of oscillation of the blood sample 14.

One of ordinary skill in the art can use other well known methods for finding the fundamental natural frequency of the blood sample throughout the hemostasis and lysis processes. For example, displacing the container 12 with a frequency function that emulates white noise having frequency components near or equal to the fundamental natural frequencies of the blood sample 14 throughout the hemostasis and lysis processes can excite the blood sample 14 to a resonant state. White noise is a frequency function that includes frequency components selected within a range of frequencies. Because the blood sample will respond with resonant excitation to a frequency that is equal or near its fundamental natural frequency, a white noise having such a frequency component will excite the blood sample 14 to a resonant state. One of ordinary skill in the art will readily appreciate that well known methods such as Fourier Frequency Analysis can be utilized to find the fundamental frequency of the blood sample 14 after being excited by white noise.

An exemplary device employing the foregoing method of determining hemostasis characteristics of a blood sample 14 will now be described. Referring to FIG. 1, the shaker 16 displaces the container 12 to excite the blood sample 14 to resonant vibration. Generally, the shaker 16 is a device capable of oscillating the container 12 with a desired frequency and amplitude. One of ordinary skill in the art will appreciate the numerous devices by which an object can be oscillated. In the disclosed example, the shaker 16 is a dipcoil, which is similar to a voice coil of a speaker. In other words, the shaker 16 includes an electromagnet that oscillates relative to a stationary permanent magnet by having its current driven by an electrical signal. The shaker 16 may be connected either directly or with a connecting link 36 to the container 12. The connecting link 36 transfers the motion created by the shaker 16 to the container 12. As is well known to those of ordinary skill in the art, characteristics of the electrical signal, i.e., voltage, current, direction of current, etc., determine the characteristics of the oscillatory motion of

the shaker 16. Accordingly, the shaker 16 can displace the container 12 with any desired amplitude and frequency within the operational limits of the shaker 16.

The container 12 holds the blood sample 14 during the excitation of the blood sample 14. The container 12 may be any shape or size. However, the shape and size of the container may affect the operation of the analyzer 10, because the container 12 acts as a resonator. The larger the container 12, the lower the natural frequency of the blood sample 14 will be. Furthermore, the container 12 cannot be too small so that a meniscus effect is produced due to the surface tension in the blood sample 14. Conversely, if the container 12 is too large, a large blood sample 14 will be needed for the analysis in the analyzer 10, which may not be medically acceptable.

An exemplary container 12 is shown in FIG. 3. The container 12 has a lower portion 40 and an upper portion 42. The lower portion 40 and the upper portion 42 are generally rectangular. The upper portion 42 has a larger width, a larger length, and a smaller depth than the lower portion 40, so as to provide an internal step 44. The container 12 also includes a lid 46 that is sealably attached to the top of the upper section 40. The container 12 includes a port 48 for receiving a blood sample 14. To reduce the meniscus effect of the blood sample 14 when placed in the container 12, the lower portion 40 is filled with the blood sample up to where the upper portion 42 begins. Accordingly the volume of the blood sample 14 is substantially equal to the volume of the lower portion 40.

To prevent the blood sample 14 from evaporating during testing and to prevent contamination thereof, the port 48 may be self sealing. For example, the port 48 may be constructed from rubber or silicon so that when a syringe needle is inserted therein, the rubber or silicon resiliently surrounds the syringe needle to substantially seal the port during the injection of the blood sample 14 into the container 12. When the needle is withdrawn from the port 48, resilience of the rubber or the silicon substantially re-seals the hole created by the needle. To prevent evaporation of the blood sample 14 and any reaction the blood sample may have by being exposed to air, the container 12 can be pre-filled or pressurized with an inert gas, such as Helium. Alternately, the air in the container can be removed to provide a vacuum inside the container 12. One of ordinary skill in the art will recognize that the pressure in the container 12 has minimal to no effect on the fundamental natural frequency of the blood sample 14. In the example disclosed herein, the container 12 is safely disposable and can be safely discarded after each use. The disposability of the container 12 ensures that the blood sample 14 is safely handled during testing and safely discarded after testing. In addition, the disposable container 12 can be manufactured to be completely sealed and only provide access thereto by the port 48. Thus, the disposability of the container 12, combined with the container 12 being completely sealed, ensure that the blood sample 14 is not exposed to air (i.e., to prevent the drying of the surface of the blood sample 14) or any other contaminants, and furthermore, ensure safety in handling and disposing of the blood sample 14 before, during, and after testing.

The analyzer 10 includes a slot (not shown) to receive the container 12. One of ordinary skill in the art will readily appreciate that the container 12 may be inserted in and removed from the slot in any manner desirable. However, to provide easy insertion and removal of the container 12 from the analyzer 10, the container 12 may include a handle (not shown) that can be held by a user for insertion and removal of the container 12 to and from the analyzer 10, respectively.

To measure oscillations of the blood sample 14 as a result of the displacement of the container 12, a fixed electromag-

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netic source **60** emits a beam **62** toward the blood sample **14**. As shown in FIG. 1, the source **60** may be part of the sensor **18** (i.e., an active sensor). Alternatively, the source **60** and a sensor **66** (i.e., a passive sensor) can be independent devices. The beam **62** is detected by the sensor **18** after being reflected from the surface of the blood sample **14**. The characteristics of the beam after being reflected from the surface of the blood sample **14** are indicative of the movement of the blood sample **14** in response to displacements of the container **12**.

One of ordinary skill in the art will appreciate that the electromagnetic beam of the source **60** may be produced by any emission within the electromagnetic spectrum so long as the beam **62** can reflect from the surface of the blood sample **14**, and the beam's characteristics after reflecting from the surface of the blood sample **14** indicate the movement of the blood sample **14**.

In the disclosed example, the source **60** is a fixed LED (Light Emitting Diode) source that directs a beam **62** towards the blood sample **14**. The beam **62** is then reflected from the surface of the blood sample **14**. Accordingly, the container **12** has an optically transparent portion so that the beam **62** and its reflection **64** can enter and exit the container **12**, respectively. In the disclosed example, the lid **46** is transparent to light. One of ordinary skill in the art will recognize that the lid **46**, although transparent, will itself reflect some of the light in the beam **62**. To reduce the reflection of light from the lid **46**, an anti-reflective coating may be applied to the lid **46**. Such anti-reflective coatings are well known to those of ordinary skill in the art as they are applied to a variety of optical devices, such as eyeglasses, telescopes, cameras, etc. Although most liquids are highly transparent to light, the surface of blood forms a highly reflective surface so that most of the beam **62** is reflected from the surface of the blood sample **14**.

Referring to FIG. 4, the displacements of the blood sample **14** relative to a rest position are shown with dashed lines **70** having an angle δ . Accordingly, the displacement of the blood sample **14** changes the angle of the reflection **64** of the beam **62** by the same angle δ . The sensor **18** intercepts the reflection **64** of the beam **62** from the surface of the blood sample **14** and produces an electric signal indicative of the displacement of the blood sample **14**. In the disclosed example, the sensor **18** includes a plurality of photo diodes that collectively detect the displacement of the reflection of the beam **64**. The outputs of the diodes are measured differentially so that peaks in the displacement of the blood sample **14**, which are indicative of resonance, can be identified.

In others example of the present disclosure, the vibrations in the blood sample **14** may be measured by a number of other devices. In one example, acoustic sensors (not shown) disposed in the container **12** can differentially measure the distance from the surface of the blood sample **14** to the sensor, which is indicative of the vibration in the blood sample **14**. In another example, electrodes (not shown) arranged in the container **12** function as either a capacitive or resistive bridge (i.e., a Wheatstone bridge). The voltage differential of the capacitors or the resistors is indicative of the vibrations of the blood sample **14**. In yet another example, two photo diodes (not shown) can be placed on an interior wall of the container near the surface of the blood sample **14**. As the blood sample **14** vibrates, it partially or fully obscures one or both of the diodes (i.e., preventing light from reaching the diodes). Accordingly the outputs of the diodes are measured differentially so that peaks in the displacement of the blood sample **14**, which are indicative of resonance, can be identified.

One of ordinary skill in the art will appreciate the numerous methods and devices that can be used for driving the shaker **16**

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and analyzing the signals from the sensor **18** for determining the hemostasis characteristics of the blood sample **14**. For instance, as shown in FIG. 5, the blood hemostasis analyzer **10** can include an internal computing device **80** that includes the necessary hardware and software to drive the shaker **16** independently or in response to signals from the sensor **18**. Furthermore, the internal computing device **80** can analyze the signals from the sensor **18** to determine the fundamental natural frequencies of the blood sample **14** during hemostasis. As described in the foregoing, such an analysis will yield data for constructing the curves **30** and other data regarding the hemostasis characteristics of the blood sample **14**. In another example as shown in FIG. 6, the analyzer **10** can include a memory device **82** for storing the data from the sensor **18** for later analysis by an external computing device **84**. The shaker **10** can be driven by a predetermined method stored in the memory device **82**, or by the external computing device **84**. In yet another example shown in FIG. 7, the analyzer **10** does not include any internal memory or computing device. During testing, the analyzer **10** is in continuous and real-time communication with an external computing device **86** (e.g., laptop, personal digital assistant, desktop computer, etc.). The external computing device **86** drives the shaker **16** and receives signals from sensor **18** to determine the hemostasis characteristics of the blood sample **14** as described in the foregoing. One of ordinary skill in the art will appreciate that numerous other well known methods, algorithms and devices can be utilized to drive the shaker **16**, independently or in response to signals from the sensor **18**, and determine blood hemostasis characteristics from the sensor signals. Furthermore, the determined blood hemostasis characteristics can be conveyed to a user by a variety of well known methods and devices, such as displaying data on a display screen, or printing the results on paper.

One of ordinary skill in the art will appreciate that the foregoing generalized device is very rugged and not easily susceptible to damage from being mishandled. The disclosed device has a very small number of moving parts or parts that are breakable. Furthermore, the simplicity of the disclosed device provides for quick replacement of a defective part when necessary.

Ambient vibrations or seismic noise near the analyzer **10** can disturb or influence the blood hemostasis analysis. Accordingly, the analyzer **10** can include a vibration filtering device onto which the analyzer **10** is mounted. In a first example as show in FIG. 8, the vibration filtering device is a hook **90**, from which the analyzer **10** is suspended by a cable **92**. In effect, the analyzer **10** is suspended from the hook **90** in a pendulum-like manner. Seismic noise or ambient vibration in a wide range of frequencies is dissipated through the hook **90** and the cable **92** prior to reaching the analyzer **10**. One of ordinary skill in the art will appreciate that any wire that is connected to the analyzer **10** for power or communication purposes can be carried by the cable **92** so as to not externally influence the motion of the analyzer **10** (e.g., hanging wires contacting other objects). In a second example as shown in FIG. 9, the seismic filtering device is a platform **100** that rests on a number of legs **102**. In effect, the platform **100** is an inverted pendulum. In application, the analyzer **10** is placed on the platform **100** so that any ambient vibration or seismic noise within a wide frequency range is dissipated through the platform **100** prior to reaching the analyzer **10**. One of ordinary skill in the art will appreciate many other ways of isolating noise, including use of vibration absorbing foams, spring suspension and the like.

Although certain apparatus constructed in accordance with the teachings of the invention have been described herein, the

scope of coverage of this patent is not limited thereto. On the contrary, this patent covers all examples of the teachings of the invention fairly falling within the scope of the appended claims either literally or under the doctrine of equivalents.

What is claimed is:

1. A method for measuring hemostasis comprising: providing a blood sample; exciting the blood sample to a resonant state by displacing the blood sample to create movement in the blood sample; observing the movement in the blood sample; periodically determining a resonant frequency of the blood sample to provide a plurality of resonant frequencies of the blood sample; and determining changes in the resonant frequencies of the blood sample over time, wherein the changes in the natural frequencies represent changes in the hemostasis characteristics of the blood sample over time.

2. A method according to claim 1, wherein displacing the blood sample to create movement comprises displacing the blood sample at a frequency, and incrementally varying the frequency until a resonant frequency of the blood sample is determined.

3. A method according to claim 1, wherein displacing the blood sample to create movement comprises displacing the blood sample incrementally varying the frequency from the resonant frequency to determine a plurality of resonant frequencies of the blood sample before, during and after hemostasis of the blood sample, wherein the plurality of the resonant frequencies are indicative of the hemostasis characteristics of the blood sample.

4. A method according to claim 1, wherein displacing the blood sample to create movement comprises displacing the blood sample with a frequency function having randomly selected frequency components selected from a range of frequencies; and determining a plurality of resonant frequencies of the blood sample before, during and after hemostasis of the blood sample; wherein the plurality of the resonant frequencies are indicative of the hemostasis characteristics of the blood sample.

5. A method according to claim 1, wherein displacing the blood sample to create movement comprises displacing the blood sample using a white noise function.

6. A method according to claim 1, wherein displacing the blood sample to create movement comprises displacing the blood sample to generate a time variant resonant response.

7. A method for measuring hemostasis comprising: placing a blood sample in a container having a portion transparent to an emission from a sensor; displacing the container with an

shaker to cause movement in the blood sample; measuring the movement in the blood sample with a sensor receiving the emission indicative of the movement of the blood sample, wherein the emission contact at least a portion of the blood sample before being received by the sensor; periodically determining a resonant frequency of the blood sample from the movement of the blood sample to provide a plurality of resonant frequencies of the blood sample; and determining changes in the resonant frequencies of the blood sample over time, wherein the changes in the natural frequencies represent changes in the hemostasis characteristics of the blood sample over time.

8. A method according to claim 7, further comprising: displacing the container at a frequency; incrementally varying the frequency until a resonant frequency of the blood sample is determined; and incrementally varying the frequency from the resonant frequency to determine a plurality of resonant frequencies of the blood sample before, during and after hemostasis of the blood sample; wherein the plurality of the resonant frequencies are indicative of the hemostasis characteristics of the blood sample.

9. A method according to claim 7, further comprising: displacing the container with a frequency function having randomly selected frequency components selected from a range of frequencies; and determining a plurality of resonant frequencies of the blood sample before, during and after hemostasis of the blood sample; wherein the plurality of the resonant frequencies are indicative of the hemostasis characteristics of the blood sample.

10. A method according to claim 7, wherein placing the blood sample in the container further comprises injecting the blood into the container through a self-sealing port on the container.

11. A method according to claim 7, wherein displacing the blood sample to create movement comprises displacing the blood sample using a white noise function.

12. A method according to claim 7, wherein displacing the blood sample to create movement comprises displacing the blood sample to generate a time variant resonant response.

13. A method according to claim 1, further comprising determining changes in the modulus of elasticity of the blood sample over time based upon the plurality of resonant frequencies, wherein changes in the modulus of elasticity represent changes in the hemostasis characteristics of the blood sample over time.

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